

= > e arachidonic acid

E1 285 ARACHIDONATE/BI
E2 83 ARACHIDONIC/BI
E3 0 --> ARACHIDONIC ACID/BI
E4 23 ARACHIDONIN/BI
E5 9 ARACHIDONO/BI
E6 1 ARACHIDONON/BI
E7 1 ARACHIDONONI/BI
E8 1 ARACHIDONONITRI/BI
E9 1 ARACHIDONONITRILE/BI
E10 1 ARACHIDONONYL/BI
E11 1 ARACHIDONONYLLECITHIN/BI
E12 52 ARACHIDONOYL/BI

= > e arachidonic acid/CN

E13 1 ARACHIDONATE-SPECIFIC PHOSPHOLIPASE A2/CN
E14 1 ARACHIDONIC 5-LIPOXYGENASE/CN
E15 1 --> ARACHIDONIC ACID/CN
E16 1 ARACHIDONIC ACID (N,2,2-3H)ETHANOLAMIDE/CN
E17 1 ARACHIDONIC ACID .OMEGA.-1 HYDROXYLASE (MOUSE STRAIN C57BL/6
J CLONE WQ2J9-7 GENE CYP2J9)/CN
E18 1 ARACHIDONIC ACID .OMEGA.-1-HYDROXYLASE/CN
E19 1 ARACHIDONIC ACID .OMEGA.-HYDROXYLASE/CN
E20 1 ARACHIDONIC ACID 12S-LIPOXYGENASE/CN
E21 1 ARACHIDONIC ACID 15-LIPOXYGENASE/CN
E22 1 ARACHIDONIC ACID 18(R)-HYDROXYLASE/CN
E23 1 ARACHIDONIC ACID 5-LIPOXYGENASE/CN
E24 1 ARACHIDONIC ACID ANHYDRIDE/CN

= > s e15

L1 1 "ARACHIDONIC ACID"/CN

= > d L1

L1 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN

RN 506-32-1 REGISTRY

CN 5,8,11,14-Eicosatetraenoic acid, (5Z,8Z,11Z,14Z)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 5,8,11,14-Eicosatetraenoic acid, (all-Z)- (8CI)

OTHER NAMES:

CN (all-Z)-5,8,11,14-Eicosatetraenoic acid

CN 5,8,11,14-all-cis-Eicosatetraenoic acid

CN 5-cis,8-cis,11-cis,14-cis-Eicosatetraenoic acid

CN 5Z,8Z,11Z,14Z-Eicosatetraenoic acid

CN all-cis-5,8,11,14-Eicosatetraenoic acid

CN arachidonate

CN Arachidonic acid

CN cis-.DELTA.5,8,11,14-Eicosatetraenoic acid

FS STEREOSEARCH

DR 10417-93-3, 929-92-0

MF C20 H32 O2

CI COM

LC STN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS,
BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB, CEN,
CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHM, CSNB, DDFU, DETHERM*,
DIOGENES, DRUGU, EMBASE, GMELIN*, HODOC*, IFICDB, IFIPAT, IFIUDB, IPA,
MEDLINE, MRCK*, NAPRALERT, NIOSHTIC, PROMT, RTECS*, SPECINFO, TOXCENTER,
USPAT2, USPATFULL, VETU, VTB

(*File contains numerically searchable property data)

Other Sources: EINECS**

(**Enter CHEMLIST File for up-to-date regulatory information)

Double bond geometry as shown.

=> e decosahecanoic acid/Cn

E25 1 DECORTISAL/CN
E26 1 DECORTISYL/CN
E27 0 --> DECOSAHECANOIC ACID/CN
E28 1 DECOSE/CN
E29 1 DECOSE, 2,7:6,10-DIANHYDRO-4,5,8,9-TETRADEOXY-3-O-(1-ETHOXYETHYL)-/CN
E30 1 DECOSERPYL/CN
E31 2 DECOSIDE/CN
E32 1 DECOSILK ART/CN
E33 1 DECOSILK BLACK OXIDE/CN
E34 1 DECOSILK DEEP BLACK/CN
E35 1 DECOSILK WHITE NY/CN
E36 1 DECOSILK WHITE NY-A 2/CN

=> e decosahecanoic acid

E37 2 DECORV/BI
E38 9 DECOS/BI
E39 0 --> DECOSAHECANOIC ACID/BI
E40 72 DECOSE/BI
E41 2 DECOSEPTAN/BI
E42 1 DECOSEPTANOSE/BI
E43 1 DECOSEPTANOSIDE/BI
E44 1 DECOSERP/BI
E45 1 DECOSERPYL/BI
E46 3 DECOSIDE/BI
E47 5 DECOSILK/BI
E48 3 DECOSOFT/BI

=> e docosahecanoic acid

E49 1 DOCOSAHECANENOATE/BI
E50 2 DOCOSAHECANOIC/BI
E51 0 --> DOCOSAHECANOIC ACID/BI
E52 7 DOCOSAHECAYN/BI
E53 14 DOCOSAHECAYNE/BI
E54 1 DOCOSAHECAYNOIC/BI
E55 6 DOCOSAHECAYNYL/BI
E56 6 DOCOSAHECAYNYLENE/BI
E57 1 DOCOSAHECEN/BI
E58 1 DOCOSAHECENO/BI
E59 1 DOCOSAHECENOATE/BI
E60 1 DOCOSAHECENOIC/BI

=> e docosahecanoic acid/cn

E61 1 DOCOSAHECAENOYL CHLORIDE, (ALL-Z)-/CN
E62 1 DOCOSAHECAENOYL COA SYNTHETASE/CN
E63 0 --> DOCOSAHECANOIC ACID/CN
E64 1 DOCOSAISOPROPOXYDECATITANOXANE/CN
E65 1 DOCOSALENE/CN
E66 1 DOCOSALENE, 1,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,
20,21,22,23,24,25,26,27,28,29,30,31,32,33,34,35,36,37,38,39,
40-TETRACONTAHYDRO-/CN
E67 1 DOCOSALENE, 1,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,

20,21,22,23,24,25,26,27,28,29,30,31,32,33,34,35,36,37,38,39,
40-TETRACONTAHYDRO-, (E)-/CN
E68 1 DOCOSALENE, 1,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,
20,21,22,23,24,25,26,27,28,29,30,31,32,33,34,35,36,37,38,39,
40-TETRACONTAHYDRO-, (Z)-/CN
E69 1 DOCOSAMETHYLCYCLOUNDECASILOXANE/CN
E70 1 DOCOSAMETHYLDECAGERMANE/CN
E71 1 DOCOSAMETHYLDECASILANE/CN
E72 1 DOCOSAMETHYLDECASILOXANE/CN

= > e docosaheptaenoic acid

E73 98 DOCOSAHEXAENOATE/BI
E74 742 DOCOSAHEXAENOIC/BI
E75 0 --> DOCOSAHEXAENOIC ACID/BI
E76 1 DOCOSAHEXAENOIN/BI
E77 21 DOCOSAHEXAENOYL/BI
E78 3 DOCOSAHEXAENOYLGLYCER/BI
E79 3 DOCOSAHEXAENOYLGLYCEROL/BI
E80 1 DOCOSAHEXAENOYLLECITHIN/BI
E81 1 DOCOSAHEXAENOYLOKADAIC/BI
E82 2 DOCOSAHEXAENOYLOXY/BI
E83 1 DOCOSAHEXAENOYLPHOSPHA/BI
E84 1 DOCOSAHEXAENOYLPHOSPHATID/BI

= > e docosaheptaenoic acid/cn

E85 1 DOCOSAHEXAENE, 1,1',1''-(1,2,3-PROPANETRIYLTRIS(OXY))TRIS-/C
N
E86 1 DOCOSAHEXAENOATE 1-MONOOXYGENASE/CN
E87 3 --> DOCOSAHEXAENOIC ACID/CN
E88 1 DOCOSAHEXAENOIC ACID ESTER WITH POLYGLYCERIN/CN
E89 1 DOCOSAHEXAENOIC ACID MONOOXYGENASE/CN
E90 1 DOCOSAHEXAENOIC ACID POLYETHYLENE GLYCOL ESTER/CN
E91 1 DOCOSAHEXAENOIC ACID, (((2,3-DIHYDROXYPROPOXY)HYDROXYPHOSPH
NYL)OXY)((1-OXOHEXADECYL)OXY)PROPYL ESTER/CN
E92 1 DOCOSAHEXAENOIC ACID, (1R)-1-(((2-AMINOETHOXY)HYDROXYPHOSPH
INYL)OXY)METHYL)-1,2-ETHANEDIYL ESTER/CN
E93 1 DOCOSAHEXAENOIC ACID, (1R)-1-(((2-AMINOETHOXY)HYDROXYPHOSPH
INYL)OXY)METHYL)-2-(((9Z)-1-OXO-9-OCTADECENYL)OXY)ETHYL ESTE
R, (Z,Z,Z,Z,Z,Z)-/CN
E94 1 DOCOSAHEXAENOIC ACID, (1R)-1-(((2-AMINOETHOXY)HYDROXYPHOSPH
INYL)OXY)METHYL)-2-((1-OXOHEXADECYL)OXY)ETHYL ESTER, (Z,Z,Z,
Z,Z,Z)-/CN
E95 1 DOCOSAHEXAENOIC ACID, (1R)-1-(((2-AMINOETHOXY)HYDROXYPHOSPH
INYL)OXY)METHYL)-2-((1-OXOHEXADECYL)OXY)ETHYL ESTER, DILITHI
UM SALT/CN
E96 1 DOCOSAHEXAENOIC ACID, (1R)-1-(((2-AMINOETHOXY)HYDROXYPHOSPH
INYL)OXY)METHYL)-2-((1-OXOOCTADECYL)OXY)ETHYL ESTER, (Z,Z,Z,
Z,Z,Z)-/CN

= > s e87

L2 3 "DOCOSAHEXAENOIC ACID"/CN

= > d l2

L2 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2004 ACS on STN
RN 32839-18-2 REGISTRY
CN Docosaheptaenoic acid, (Z,Z,Z,Z,Z,Z)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:

STN SEARCH PERFORMED ON 01/30/2004

CN Docosahexaenoic acid, (all-Z)- (8CI)

OTHER NAMES:

CN cis-Docosahexaenoic acid

CN Docosahexaenoic acid

DR 179092-16-1

MF C22 H32 O2

CI IDS, COM

LC STN Files: ADISNEWS, AGRICOLA, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CEN, CIN, EMBASE, PROMT, TOXCENTER, USPAT2, USPATFULL

CM 1

CRN 112-85-6

CMF C22 H44 O2

1436 REFERENCES IN FILE CA (1907 TO DATE)

35 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

1436 REFERENCES IN FILE CAPLUS (1907 TO DATE)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU, DGENE, DRUGB, DRUGMONOG2, ...' ENTERED AT 14:49:27 ON 30 JAN 2004

= > (L1 AND L2) AND (PARTICLE OR MATTER OR FORMULA? OR COMPOSITION OR MATTER OR MIXTURE)

0* FILE ADISCTI

2 FILE ADISNEWS

150 FILE AGRICOLA

7 FILE ANABSTR

0* FILE AQUASCI

31 FILE BIOBUSINESS

0* FILE BIOCOMMERCE

485 FILE BIOSIS

50 FILE BIOTECHNO

0* FILE CABA

0* FILE CAPLUS

0* FILE CEABA-VTB

1 FILE CEN

14 FILE CIN

0* FILE CONFSCI

19 FILES SEARCHED...

0* FILE CROPB

0* FILE CROPU

0* FILE DISSABS

0* FILE DDFB

0* FILE DDFU

0* FILE DGENE

0* FILE DRUGB

0* FILE DRUGU

29 FILES SEARCHED...

0* FILE EMBAL

0* FILE ESBIODASE

0* FILE FEDRIP

0* FILE FOMAD

0* FILE FOREGE

0* FILE FROSTI

0* FILE GENBANK

0* FILE HEALSAFE

0* FILE IFIPAT

0* FILE KOSMET

0* FILE LIFESCI

0* FILE MEDICONF
 0* FILE NTIS
 0* FILE NUTRACEUT

50 FILES SEARCHED...

0* FILE OCEAN
 0* FILE PASCAL
 0* FILE PCTGEN
 0* FILE PHARMAML
 0* FILE PHIC
 0* FILE PHIN
 13 FILE PROMT
 0* FILE RDISCLOSURE
 0* FILE SCISEARCH
 56 FILE TOXCENTER
 0* FILE USPATFULL

63 FILES SEARCHED...

0* FILE USPAT2
 0* FILE VETB
 0* FILE VETU

67 FILES SEARCHED...

L3 QUE (L1 AND L2) AND (PARTICLE OR MATTER OR FORMULA? OR COMPOSITION OR MATTER OR MIXTURE) 10 FILES HAVE ONE OR MORE ANSWERS

L4 QUE MICROORGANISM OR FUNG? OR BCTERIA OR DINOFLAGELLATE OR ALGAE OR MORTIERELLA OR CRYPTHECODINIUM 68 FILES HAVE ONE OR MORE ANSWERS

L5 QUE (PHOSPHOLIPID) AND (L1 AND L2) 7 FILES HAVE ONE OR MORE ANSWERS

L6 QUE (L1 AND L2) AND L4 8 FILES HAVE ONE OR MORE ANSWERS

L7 QUE L5 AND L6 4 FILES HAVE ONE OR MORE ANSWERS

L8 QUE L3 AND L7 , 1 FILES HAVE ONE OR MORE ANSWERS

=> D RANK

F1 5 BIOSIS

L9 5 L3 AND L7

L10 5 DUP REM L9 (0 DUPLICATES REMOVED)

L10 ANSWER 1 OF 5 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

AB Objective: To review briefly the influence of dietary long-chain polyunsaturated fatty acids (LC-PUFA) on tissue composition and functionality in early infancy. Moreover, the influences of LC-PUFA sources on plasma composition as well as the effects of these fatty acids on intestinal repair after malnutrition are discussed.
 Results: Human milk not only supplies essential fatty acids but also contains up to 2% of the total fatty acids as LC-PUFA, of which arachidonic acid (AA) and docosahexaenoic acid (DHA) are considered the most important. Plasma and erythrocyte levels of both AA and DHA are decreased in infants fed artificial standard milk formulae. However, the supplementation of formulae with these fatty acids in amounts close to that of human milk leads to tissue LC-PUFA patterns similar to those of breastfed infants. However, the bioavailability of LC-PUFA depends on the typical LC-PUFA source; egg phospholipids increases both AA and DHA in plasma phospholipids and HDL more than a mixture of tuna and fungal triglycerides.
 Conclusions: Dietary LC-PUFA affects positively the growth and development of the infant and ameliorates the visual and cognitive functions, particularly in preterm infants. Likewise, LC-PUFA improves intestinal repair in severe protein-energy malnutrition; therefore, its qualitative and quantitative dietary supply should be considered.

AN 2003:540636 BIOSIS

DN PREV200300543231

TI Role of long-chain polyunsaturated fatty acids in infant nutrition.
AU Gil, A. [Reprint Author]; Ramirez, M.; Gil, M.
CS Department of Biochemistry and Molecular Biology, School of Pharmacy,
University of Granada, Campus Universitario de Cartuja, 18071, Granada,
Spain
agil@ugr.es
SO European Journal of Clinical Nutrition, (September 2003) Vol. 57, No.
Supplement 1, pp. S31-S34. print.
CODEN: EJCNEQ. ISSN: 0954-3007.
DT Article
General Review; (Literature Review)
LA English
ED Entered STN: 19 Nov 2003
Last Updated on STN: 19 Nov 2003

L10 ANSWER 2 OF 5 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

AB Addition of arachidonic acid (AA) and docosahexaenoic acid (DHA) to infant formula promotes visual and neural development. This study was designed to determine whether the source of dietary long-chain polyunsaturated fatty acids (LCPUFA) affected overall animal health and safety. Piglets consumed ad libitum from 1 to 16 d of age a skim milk-based formula with different fat sources added to provide 50% of the metabolizable energy. Treatment groups were as follows: control (CNTL; no added LCPUFA), egg phospholipid (PL), algal/fungal triglyceride (TG) oils, TG plus PL (soy lecithin source) added to match phospholipid treatment (TG + PL) and essential fatty acid deficient (EFAD). Formulas with LCPUFA provided 0.6 and 0.3 g/100 g total fatty acids as AA and DHA, respectively. CNTL piglets had 40% longer ileal villi than PL piglets ($P < 0.03$), but the TG group was not different from the CNTL group. Gross liver histology did not differ among any of the formula-fed groups ($P > 0.1$). Apparent dry matter digestibility was 10% greater in CNTL, TG and TG + PL groups compared with PL piglets ($P < 0.002$). No differences in alanine aminotransferase were detected among treatments, but aspartate aminotransferase was elevated ($P < 0.03$) in PL piglets compared with TG + PL piglets. Total plasma AA concentration was greater in the TG group compared with CNTL piglets ($P < 0.05$). Total plasma DHA concentrations were greater in TG piglets compared with PL ($P < 0.06$) or CNTL ($P < 0.02$) piglets. These data demonstrate that the algal/fungal TG sources of DHA and AA may be a more appropriate supplement for infant formulas than the egg PL source based on piglet plasma fatty acid profiles and apparent dry matter digestibilities.

AN 2002:588302 BIOSIS

DN PREV200200588302

TI Comparison of triglycerides and phospholipids as supplemental sources of dietary long-chain polyunsaturated fatty acids in piglets.
AU Mathews, Susan A.; Oliver, William T.; Phillips, Oulayvanh T.; Odle, Jack; Diersen-Schade, Deborah A.; Harrell, Robert J. [Reprint author]
CS North Carolina State University, Raleigh, NC, 27695, USA
bob_harrell@ncsu.edu
SO Journal of Nutrition, (October, 2002) Vol. 132, No. 10, pp. 3081-3089.
print.
CODEN: JONUAI. ISSN: 0022-3166.
DT Article
LA English
ED Entered STN: 13 Nov 2002
Last Updated on STN: 13 Nov 2002

L10 ANSWER 3 OF 5 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

AN 2000:446603 BIOSIS

DN PREV200000446603

TI Dietary long-chain polyunsaturated fatty acids in the form of phospholipids or triglycerides influence plasma lipoproteins composition.

AU Amate, L.; Gil, A. [Reprint author]; Ramirez, M.

CS Biochemistry and Molecular Biology Dpt, University of Granada, Granada, Spain

SO Clinical Nutrition (Edinburgh), (August, 2000) Vol. 19, No. Supplement 1, pp. 18. print.

Meeting Info.: 22nd Congress of the European Society of Parenteral and Enteral Nutrition. Madrid, Spain. September 09-13, 2000.

CODEN: CLNUDP. ISSN: 0261-5614.

DT Conference; (Meeting)

Conference; Abstract; (Meeting Abstract)

LA English

ED Entered STN: 18 Oct 2000

Last Updated on STN: 10 Jan 2002

L10 ANSWER 4 OF 5 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

AB Critically ill hospital patients were fed enteral formulas containing different fat substrates. Seven patients received formula X, which contained 28 g of structured triglycerides and menhaden oil to provide 7.6 g of medium-chain fatty acids, 2.5 g linoleic acid, 1.3 g eicosapentaenoic acid, and 0.4 g docosahexaenoic acid per 1000 mL of formula. Six patients received formula Y consisting of 36.8 g of medium-chain triglycerides and corn and soy oils providing 14.3 g medium-chain fatty acids and 11.7 g linoleic acid per 1000 mL. Feeding of formula X increased plasma total phospholipid levels of eicosapentaenoic acid on days 7 and 14 and docosahexaenoic acid levels on day 14. Plasma levels of linoleic acid were reduced in formula-X-fed in comparison to formula -Y-fed patients, whereas arachidonic acid was maintained in both groups during feeding. As a result of these changes, the patients receiving formula X had decreased ratios of arachidonic acid:eicosapentaenoic acid in plasma. Formula Y feeding did not alter eicosapentaenoic acid and docosahexaenoic acid levels in the plasma. In the erythrocyte, formula X feeding resulted in a threefold increase in eicosapentaenoic acid from mean baseline levels of 0.4 \pm 0.4% to a mean value of 1.2 \pm 0.9% at day 7. The formula X feeding decreased linoleic acid levels on days 7 and 14, whereas levels of arachidonic acid and docosahexaenoic acid remained constant. Formula Y feeding did not affect any of the parameters measured for erythrocytes. The ability to alter plasma and erythrocyte levels of n-3 fatty acids and plasma arachidonic acid:eicosapentaenoic acid ratios may have important implications for patients at risk for sepsis.

AN 1993:167617 BIOSIS

DN PREV199395088667

TI Changes in plasma and erythrocyte fatty acids in patients fed enteral formulas containing different fats.

AU Adams, Steve; Yeh, Yu-Yan; Jensen, Gordon L. [Reprint author]

CS Geisinger Med. Center, Danville, PA 17822, USA

SO Journal of Parenteral and Enteral Nutrition, (1993) Vol. 17, No. 1, pp. 30-34.

CODEN: JPENDU. ISSN: 0148-6071.

DT Article

LA English

ED Entered STN: 31 Mar 1993

Last Updated on STN: 1 Apr 1993

L10 ANSWER 5 OF 5 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

AB The phospholipid and fatty acid compositions of the host infected erythrocyte plasma membrane (IEPM) have been determined for erythrocytes infected with the human malaria parasite *Plasmodium falciparum*. IEPM were prepared by selective lysis of the host erythrocyte (but not of the parasite membranes) with 0.1% saponin, followed by differential centrifugation. The purity of the IEPM was determined by measuring the membrane-specific enzyme markers acetylcholinesterase, glutamate dehydrogenase and lactate dehydrogenase, and by immunoelectron microscopy using monoclonal antibodies specific for human erythrocyte glycophorin A (4E7) and for a 195 kDa parasite membrane glycoprotein (Pf6 3B10.1). Both approaches demonstrated that the host erythrocyte plasma

membrane preparation was free from contamination by parasite membranes. During intra-erythrocytic development of the parasite, the phospholipid composition of the erythrocyte membrane was strikingly altered. IEPM contained more phosphatidylcholine (38.7% versus 31.7%) and phosphatidylinositol (2.1% versus 0.8%) and less sphingomyelin (14.6% versus 28.0%) than normal uninfected erythrocytes. Similar alterations in phospholipid composition were determined for erythrocyte membranes of parasitized cells isolated by an alternative method utilizing polycationic polyacrylamide microbeads (Affigel 731). The total fatty acid compositions of the major phospholipids in IEPM were determined by g.l.c. The percent of polyunsaturated fatty acids in normal erythrocyte phospholipids (39.4%) was much higher than in phospholipids from purified parasites (23.3%) of IEPM (24.0%). The unsaturation index of phospholipids in IEPM was considerably lower than in uninfected erythrocytes (107.5 versus 161.0) and was very similar to that in purified parasites (107.5 versus 98.5). Large increases in palmitic acid (C16:0) (from 21.88% to 31.21%) and in oleic acid (C18:1) (from 14.64% to 24.60%), and major decreases in arachidonic acid (C20:4) (from 17.36% to 7.85%) and in docosahexaenoic acid (C22:6) (from 4.34% to 1.8%) occurred as a result of infection. The fatty acid profiles of individual phospholipid classes from IEPM resembled in many instances the fatty acid profiles of parasite phospholipids rather than those of uninfected erythrocytes. Analysis of IEPM from *P. falciparum*-infected erythrocytes (trophozoite stage) revealed that, during intra-erythrocytic maturation of the parasite, the host erythrocyte phospholipid composition was markedly refashioned. These alterations were not dependent on the method used to isolate the IEPM, with similar results obtained using either a saponin-lysis method or binding to Affigel beads. Since mature erythrocytes have negligible lipid synthesis and metabolism, these alterations must occur as a result of parasite-directed metabolism of erythrocyte lipids and/or trafficking of lipids between the parasite and erythrocyte membranes. AN 1991:230569 BIOSIS DN PREV199191122029; BA91:122029T1 MODIFICATION OF HOST CELL MEMBRANE LIPID COMPOSITION BY THE INTRA-ERYTHROCYTIC HUMAN MALARIA PARASITE *PLASMODIUM FALCIPARUM*.

AU HSIAO L L [Reprint author]; HOWARD R J; AIKAWA M; TARASCHI T F

CS DEP PATHOLOGY CELL BIOL, THOMAS JEFFERSON UNIV, 1020 LOCUST ST, PHILADELPHIA, PA 19107, USA

SO Biochemical Journal, (1991) Vol. 274, No. 1, pp. 121-132.

ISSN: 0264-6021.

DT Article

FS BA

LA ENGLISH

ED Entered STN: 9 May 1991

Last Updated on STN: 16 Jul 1991